CRF Problem Report



The Scientific and Technical Information Center (STIC) experienced problem when processing the following computer readable form (CRF):

7 9 2001 Application Serial Number 09/772 134

Filing Date:

Date Processed by STIC: 2-9-01

STIC Contact: Mark Spencer, 703-308-4212

Nature of Problem:

The CRF (was):

(circle one) Damaged or Unreadable (for Unreadable, see attached)

Blank (no files on CRF) (see attached)

Empty file (filename present, but no bytes in file) (see attached)

Virus-infected. Virus name: _____ The STIC will not process the CRF.

Not saved in ASCII text

Sequence Listing was embedded in the file. According to Sequence Rules, submitted file should only be the Sequence Listing.

Did not contain a Sequence Listing. (see attached sample)

V Other: Non-valid format, Example of correct formet attached

PLEASE USE THE CHECKER VERSION 3.0 PROGRAM TO REDUCE ERRORS. SEE BELOW FOR DETAILS:

Checker Version 3.0

The Checker Version 3.0 application is a state-of the-art Windows based software program employing a logical and intuitive user-interface to check whether a sequence listing is in compliance with format and content rules. Checker Version 3.0 works for sequence listings generated for the original version of 37 CFR §§1.821 - 1.825 effective October 1, 1990 (old rules) and the revised version (new rules) effective July 1, 1998 as well as World Intellectual Property Organization (WIPO) Standard ST.25.

Checker Version 3.0 replaces the previous DOS-based version of Checker, and is Y2Kcompliant. Checker allows public users to check sequence listings in Computer Readable form (CRF) before submitting them to the United States Patent and Trademark Office (USPTO). Use of Checker prior to filing the sequence listing is expected to result in fewer errored sequence listings, thus saving time and money.

Checker Version 3.0 can be down loaded from the USPTO website at the following address: http://www.uspto.gov/web/offices/pac/checker

Appendix A To Subpart C to Part 1-Sample Sequence Listing

OT 2 9 2001 A

<110> Smith, John

Smith, Jane

<120> Example of a Sequence Listing

<130> 01-00001

<140> US 08/999,999

<141> 1998-02-28

<150> EP 91000000

<151> 1997-12-31

<170> PatentIn ver. 2.0

<210> 1

<211> 403

<212> DNA

<213> Paramecium aurelia

<220>

<221> CDS

<222> 341..394

<300>

<301> Doe, Richard

<302> Isolation and Characterization of a Gene Encoding a

Protease from Paramecium sp.

<303> Journal of Fictional Genes

<305> 4

<306> 1 - 7

<307> 1988-06-20

<400> 1

ctactctact ctactctcat ctactatett ctttggatet ctgagtetge ctgagtggta

ctcttgagtc ctggagatct ctcctctcac atgtgatcgt cgagactgac cgatagatcg 120

ctgactgact ctgagatagt cgagcccgta cgagacccgt cgagggtgac agagagtggg 180

cgcgtgcgcg cagagcgccg cgccggtgcg cgcgcgagtg cgcggtgggc cgcgcgaggg 240

ctttcgcggc agcggcggcg ctttccggcg cgcgcccgtc cgcccctaga cctgagaggt

cttctcttcc ctcctctca ctagagaggt ctatatatac atg gtt tca atg ttc

Met Val Ser Met Phe

age ttg tet tte aaa tgg eet gga ttt tgt ttg ttt gtt tgtttgete

403

Ser Leu Ser Phe Lys Trp Pro Gly Phe Cys Leu Phe Val

10

15

<210> 2

<211> 18

<212> PRT

<213> Paramecium aurelia

<400> 2

bl=630No.;104/Monday, June 1, 199

Met Val Ser Met Phe Ser Leu Ser Phe Lys Trp Pro Gly Phe dys Leu

5

10

Phe Val

1

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<130>	•••••••	File Reference		Personal file reference		M. M when filed prior to assignment of appl. nu. ber.
<140>		Comment	l		••••••	ber. ber. ber.
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: '<151>	••••••	Prior Application Filin	19 🕸	Specify as: yyyy-mm-dd		00.000.119.410.120
<160>		Date. Number of SEQ ID N		·		M. If applicable.
<170>		Software		Count includes total number of SEQ ID NOs		M
	- 1		····· / '	Name of software used to create the Sequen	nce	O
≪10>		SEQ ID NO:1:	F	Response shall be an integer representing t		
. <2115	ı	~ .		SEQ-ID NO shown.	ine	M.
1		Length	F	Respond with an integer expression the numb	per l	M. • • •
TO TO COLOR			- 	of bases or amino acid residues.		
Numeric lo	en-	Definition		Comments and format		
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<212>		Туре	w	Whether presented sequence molecule is DN/		4.
			- 1	RNA, or PRT (protein). If a nucleotide so	c.	To a secondary
	1		- 1	quence contains both DNA and RNA frac	n-	,
	- 1			ments, the type shall be "DNA." In addition the combined DNA/RNA molecule shall be full	n.	A.
	- 1		-	ther described in the <220> to <223> feature	(-	
-213s	1	Dennais	- 1 - 1	section.	- 1	
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				Artificial Sequence. In addition, the "Unknown or "Artificial Sequence" organisms shall be fur	.	gara a di
	- 1		'i	ther described in the <220> to <223> feature		
<220s	ے ا	eature	5	section,	- 1	***************************************
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	- 1		"	" we sequence		ICO DOSC WAS USED IN A RECUIRCOCK IF COCK
<221>		ama/Va	_		;	VISM is "Artificial Sequence" or "Unknown'; if molecule is combined DNA/RNA"
Q.C.17	```	ame/key	· Pro	vide appropriate Identifier for feature, pref-	M.	under the following conditions: if "a " "vas "
			6	erably from WIPO Standard ST.25 (1998), Appendix 2, Tables 5 and 6.	0	n a modified of anastral Leaming agid of meet:
<222>	Lo	cation	. Spc	cily location within sequence; where appro-	,	co pase was used in a commond
	-) pr	nate state number of first and last bases/	, ,	under the following conditions: if "n," "Xaa," r a modified or unusual L-amino acid or modi-
<223>	. 01	her Information	an	mino acids in feature.	,	ed base was used in a sequence
			0	er relevant information; four lines maximum	M.	under the following conditions: if "a " "vaa "
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					1 14	ed base was used in a sequence; if ORGA- ISM is "Antificial Sequence" or "Unknown"; if
<300>	Pul	blication Information	Leav	ve blank after <300>	• • • • • • • • • • • • • • • • • • • •	olecule is combined DNA/RNA.
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<305> <306>	Issu	e			O. O.	i
<307>	Date	es			Ö.	·
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<313>	Relev	rant Residuesz	MOH	(position) TO (position)	,	
<400>	Sequ		SEC II	D NO should follow the numeric identifier M		
1			ang .	should appear on the line preceding the		
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